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9		BEFORE THE BOARD OF REGISTERED NURSING					
	DEPARTMENT OF CONSUMER AFFAIRS						
10		STATE OF C	.ALIFORNI	A			
11	In the Matter of the	Accusation Against:	Case No.	2010-662			
12	CHRISTINE MAI	_					
13	4956 Narranganse	tt Avenue	ACCUS	ATION			
14	San Diego, CA 92		ACCUS	ATION			
15	Registered Nurse	License No. 748793					
16		Respondent.					
17			J.				
		-11					
18	Complainant		ATTIC .				
19			RTIES				
20	1. Louise	1. Louise R. Bailey, M.Ed., RN (Complainant) brings this Accusation solely in her					
21	official capacity as	ficial capacity as the Interim Executive Officer of the Board of Registered Nursing, Department					
22	of Consumer Affair	of Consumer Affairs.					
23	2. On or about April 9, 2009, the Board of Registered Nursing issued Registered Nurse						
24	License Number 748793 to Christine Marie Nowicki (Respondent). The Registered Nurse						
25	License was in full force and effect at all times relevant to the charges brought herein and will						
26	expire on August 3	expire on August 31, 2010, unless renewed.					
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28	111						
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JURISDICTION

- 3. This Accusation is brought before the Board of Registered Nursing (Board),
 Department of Consumer Affairs, under the authority of the following laws. All section
 references are to the Business and Professions Code unless otherwise indicated.
- 4. Section 2750 of the Business and Professions Code ("Code") provides, in pertinent part, that the Board may discipline any licensee, including a licensee holding a temporary or an inactive license, for any reason provided in Article 3 (commencing with section 2750) of the Nursing Practice Act.
- 5. Section 2764 of the Code provides, in pertinent part, that the expiration of a license shall not deprive the Board of jurisdiction to proceed with a disciplinary proceeding against the licensee or to render a decision imposing discipline on the license.
- 6. Section 2811(b) of the Code provides, in pertinent part, that each license not renewed shall expire but may within a period of eight years thereafter be reinstated upon payment of the biennial renewal fee and penalty fee and upon submission of such proof of the applicant's qualifications as may be required by the board, except that during such eight-year period no examination shall be required as a condition for the reinstatement of any such expired license which has lapsed solely by reason of non-payment of the renewal fee.

STATUTORY PROVISIONS

- 7. Section 2761, subdivision (a), states that the Board may take disciplinary action against a certified or licensed nurse or deny an application for a certificate or license for unprofessional conduct.
 - 8. Section 2762 of the Code states:
- "In addition to other acts constituting unprofessional conduct within the meaning of this chapter [the Nursing Practice Act], it is unprofessional conduct for a person licensed under this chapter to do any of the following:
- "(a) Obtain or possess in violation of law, or prescribe, or except as directed by a licensed physician and surgeon, dentist, or podiatrist administer to himself or herself, or furnish or administer to another, any controlled substance as defined in Division 10 (commencing with

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Section 11000) of the Health and Safety Code or any dangerous drug or dangerous device as defined in Section 4022.

- "(e) Falsify, or make grossly incorrect, grossly inconsistent, or unintelligible entries in any hospital, patient, or other record pertaining to the substances described in subdivision (a) of this section."
 - 9. Code section 4060 states, in pertinent part:
- "No person shall possess any controlled substance, except that furnished to a person upon the prescription of a physician, dentist, podiatrist, optometrist, veterinarian, or naturopathic doctor pursuant to Section 3640.7, or furnished pursuant to a drug order issued by a certified nurse-midwife pursuant to Section 2746.51, a nurse practitioner pursuant to Section 2836.1, a physician assistant pursuant to Section 3502.1, a naturophatic doctor pursuant to Section 3640.5, or a pharmacist pursuant to either subparagraph (D) of paragraph (4) of, or clause (iv) of subparagraph (A) of paragraph (5) of, subdivision (a) of Section 4052. This section shall not apply to the possession of any controlled substance by a manufacturer, wholesaler, pharmacy, pharmacist, podiatrist, dentist, optometrist, veterinarian, naturophatic doctor, certified nurse-midwife, nurse practitioner, or physician assistant, when in stock in containers correctly labeled with the name and address of the supplier or producer...
- 10. Health and Safety Code section 11170 states that no person shall prescribe, administer, or furnish a controlled substance for himself.
- 11. Health and Safety Code section 11173, subdivision (a) states, in pertinent part, that "[n]o person shall obtain or attempt to obtain controlled substances, or procure or attempt to procure the administration of or prescription for controlled substances, (1) by fraud, deceit, misrepresentation, or subterfuge . . ."

12. Section 125.3 of the Code provides, in pertinent part, that the Board may request the administrative law judge to direct a licentiate found to have committed a violation or violations of the licensing act to pay a sum not to exceed the reasonable costs of the investigation and enforcement of the case.

DRUGS

- 13. Hydrocodone bitartate/acetaminophen, also known by the brand names Vicodin, Norco, Zydone, Maxidone, Lortab, Lorcet, Hydrocet, Co-Gesic, and Anexsia, is a narcotic Schedule III controlled substance as designated by Health and Safety Code section 11056(e)(4), and is a dangerous drug pursuant to Business and Professions Code section 4022. Hydrocodone is used as a narcotic analgesic in the relief of pain.
- 14. Lorazepam, sold under the brand name Ativan, is a Schedule IV controlled substance as designated by Health and Safety Code section 11057(d)(16), and is a dangerous drug pursuant to Business and Professions Code section 4022. Lorazepam is used in the treatment of anxiety disorders and for short-term (up to 4 months) relief of the symptoms of anxiety.

FACTUAL ALLEGATIONS

15. Respondent was employed as a traveling registered nurse by AMN HealthCare from November 2008 until March 2009, assigned to Palomar Medical Center (PMC), Escondido, California. On or about March 5, 2009, during a routine Pyxis¹ User Audit at PMC, it was noted that Respondent was withdrawing more Vicodin and Ativan than her co-workers. A review of patient records and the Pyxis Activities for 11 shifts worked by Respondent during the month of February 2009 at PMC, revealed that Respondent made inaccurate entries in hospital and patient

Pyxis" is a trade name for the automatic single-unit dose medication dispensing system that records information such as patient name, physician orders, date and time medication was withdrawn, and the name of the licensed individual who withdrew and administered the medication. Each user/operator is given a user identification code to operate the control panel. Sometimes only portions of the withdrawn narcotics are given to the patient. The portions not given to the patient are referred to as "wastage." This waste must be witnessed by another authorized user and is also recorded by the Pyxis machine.

medical records, charted that she administered narcotic medications to patients with "0" pain levels, failed to assess and chart the pain levels of patients receiving pain medication, charted the administering of medications before their removal from Pyxis, charted the administering of medications more frequently than the 4 to 6 hours as ordered by the physician(s), and withdrew from Pyxis Hydrocodone and Lorazepam tablets that Respondent did not chart and was unable to account for, as follows:

16. Patient FIN 9016857:

- a. Physician's order(s) for this patient were as follows: Lorazepam 0.5mg every 4 hours as needed.
- b. On February 3, 2009, at 0931 hours, Respondent withdrew from Pyxis one (1) 0.5mg tablet Lorazepam and charted on the MAR that she administered the medication at 0945 hours. The medication was charted as having been given by IV versus a tablet by mouth.
- c. On February 3, 2009, at 1208 hours, Respondent withdrew from Pyxis one (1) 0.5mg tablet Lorazepam and charted on the MAR that she administered the medication at 1207 hours, 2 hours and 22 minutes after the first dose, not every 4 hours per the physician's orders.
- d. On February 3, 2009, at 1626 hours, Respondent withdrew from Pyxis one (1)
 0.5 mg. tablet Lorazepam and charted on the MAR that she administered the medication at 1600 hours, 26 minutes before the medication was removed from Pyxis.
- c. Summary: Respondent removed three (3) 0.5 mg. tablets of Lorazepam, three (3) 0.5 mg tablets were charted as given, and one (1) tablet is unaccounted for.

17. Patient FIN 9013540:

- a. The physician's order(s) for this patient were as follows: Hydrocodone 5-500mg 1-2 tablets every 4 hours as needed.
- b. On February 3, 2009, at 0802 hours, Respondent withdrew from Pyxis two (2) 5/500mg tablets of Hydrocodone and charted on the MAR that the medication was administered at 0830 hours.

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- c. On February 3, 2009, at 1230 hours, Respondent withdrew from Pyxis two (2) 5/500 tablets of Hydrocodone and charted on the MAR that the medication was administered on the same date at 1200 hours. Pain assessment of "0" was charted 30 minutes before removal.
- d. On February 3, 2009, at 1357 hours, Respondent wasted two (2) 5/500 tablets of Hydrocodone.
- e. On February 3, 2009, at 1358 hours, Respondent withdrew two (2) 5/500 mg and charted on the MAR that the medication was administered at 1650 hours. This medication was removed one minute after waste detailed at paragraph 16(d) and administered approximately three (3) hours after withdrawal, while pain assessment of "0" was charted.
- f. Summary: Respondent removed six (6) 5/500 tablets of Hydrocodone, charted that she administered six (6), and charted the wasting of two (2) tablets. Respondent charted administering two (2) tablets thirty minutes before she removed them (see paragraph 16(c)).

18. Patient FIN 9018413:

- a. The physician's order(s) for this patient were as follows: Lorazepam 4mg every six (6) hours as needed for anxiety.
- b. On February 3, 2009, at 1037 hours, Respondent withdrew from Pyxis one (1) 5/500mg tablets of Hydrocodone and charted on the MAR that the medication was administered at 1030 hours, seven minutes prior to the removal of the medication from Pyxis at 0830 hours.

19. Patient FIN 9018013:

- a. The physician's order(s) for this patient were as follows: Hydrocodone 5-500mg 1 tablet every 4 hours as needed.
- b. On February 5, 2009, at 0738 hours, Respondent withdrew from Pyxis two (2) 5/500mg tablets of Hydrocodone and charted on the MAR that the medication was administered at 0800 hours. There was no pain assessment charted, and two (2) tablets were withdrawn instead of one (1) tablet, per the physician's orders.
- c. On February 5, 2009, at 1114 hours, Respondent withdrew from Pyxis two (2) 5/500mg tablets of Hydrocodone and charted on the MAR that the medication was administered at 1200 hours. There was no pain assessment charted, two (2) tablets were withdrawn instead of

one (1) tablet, per the physician's orders, and the medication was administered 45 minutes after removal.

- d. On February 5, 2009, at 1633 hours, Respondent withdrew from Pyxis two 2) 5/500mg tablets of Hydrocodone and charted on the MAR that the medication was administered at 1700 hours. There was no pain assessment charted, two (2) tablets were withdrawn instead of one (1) tablet, per the physician's orders.
- e. On February 6, 2009, at 0825 hours, Respondent withdrew from Pyxis two (2) 5/500mg tablets of Hydrocodone and charted on the MAR that the medication was administered at 0745 hours. There was a pain assessment given 40 minutes prior to removal. Two (2) tablets were withdrawn instead of one (1) tablet, per the physician's orders.
- f. On February 6, 2009, at 1057 hours, Respondent withdrew from Pyxis two (2) 5/500mg tablets of Hydrocodone and charted on the MAR that the medication was administered at 1100 hours. There was no pain assessment charted prior to the removal of the medication; it was charted as given three (3) minutes after the removal. Two (2) tablets were withdrawn instead of one (1) tablet, per the physician's orders, and charted as given more frequently than every four (4) hours per physician's orders.
- g. On February 6, 2009, at 1502 hours, Respondent withdrew from Pyxis two (2) 5/500mg tablets of Hydrocodone and charted on the MAR that the medication was administered at 1536 hours. There was no pain assessment charted, and two (2) tablets were withdrawn instead of one (1) tablet, per the physician's orders.
- h. On February 6, 2009, at 1752 hours, Respondent withdrew from Pyxis two (2) 5/500mg tablets of Hydrocodone and not charted on the MAR as having been administered to the patient. There was no pain assessment charted, and two (2) tablets were withdrawn instead of one (1) tablet, per the physician's orders. There are two (2) 5/500mg Hydrocodone tablets unaccounted for.
- g. On February 16, 2009, at 0914 hours, Respondent withdrew from Pyxis two (2) 5/500mg tablets of Hydrocodone and charted on the MAR that the medication was administered at 0800 hours. There was no pain assessment charted, two (2) tablets were

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withdrawn instead of one (1) tablet, per the Physician's Orders, and the medication was charted as being administered 74 minutes before removal.

- h. On February 16, 2009, at 1201 hours, Respondent withdrew from Pyxis two (2) 5/500mg tablets of Hydrocodone and not charted as given. There was no pain assessment charted, and two (2) tablets were withdrawn instead of one (1) tablet per the physician's orders. Two (2) tablets of Hydrocodone are unaccounted for.
- i. Summary: On February 5, 2009, Respondent removed six (6) 5/500mg. tablets of Hydrocodone; six (6) were charted as given with no pain assessment charted. On February 6, 2009, Respondent removed eight (8) 5/500mg tablets of Hydrocodone, six (6) charted as given with no pain assessment and two (2) tablets unaccounted for. On February 16, 2009, Respondent removed four (4) 5/500mg. tablets of Hydrocodone, two (2) were charted as given with no pain assessment charted and two tablets are unaccounted for.

20. Patient FIN 8981168:

- a. The physician's order(s) for this patient were as follows: Vicodin 1-2 5-500mg tablets every four (4) hours as needed.
- b. On February 5, 2009, at 0904 hours, Respondent withdrew from Pyxis two (2) 5/500mg tablets of Hydrocodone and charted on the MAR that the medication was administered at 0946 hours. There was no pain assessment charted, and the medication was charted as administered 42 minutes after removal.
- c. On February 5, 2009, at 1151 hours, Respondent withdrew from Pyxis two (2) 5/500mg tablets of Hydrocodone and charted on the MAR that the medication was administered at 1230 hours. The pain assessment was charted as "2" and the medication was charted as given 39 minutes after removal.
- d. On February 5, 2009, at 1704 hours, Respondent withdrew from Pyxis two (2) 5/500mg tablets of Hydrocodone and charted on the MAR that the medication was administered at 1748 hours. There was no pain assessment charted and the medication was charted as given 44 minutes after removal.

e. Summary: Pyxis Patient Activity Report indicates one (1) tablet was removed on February 3, 2009 at 0258 hours, one (1) tablet on February 4, 2009, at 1202 hours, and one (1) tablet on February 5, 2009, at 0407 hours. On February 5, 2009, Respondent removed six (6) tablets, and six (6) were charted as given without a pain assessment.

21. Patient FIN 9019895:

- a. The physician's order(s) for this patient were as follows: Vicodin 5-500mg 1-2 tablets every 4 hours as needed for moderate to severe pain (4-10 of 10).
- b. On February 6, 2009, at 0805 hours, Respondent withdrew from Pyxis two (2) 5/500mg tablets of Hydrocodone and charted on the MAR that the medication was administered at 0853 hours. The pain assessment was charted as "4" and the medication was charted as having been administered 48 minutes after removal.
- c. On February 6, 2009, at 1233 hours, Respondent withdrew from Pyxis two (2) 5/500mg tablets of Hydrocodone and charted on the MAR that the medication was administered at 1245 hours. The pain assessment was not charted.
- d. On February 6, 2009, at 1619 hours, Respondent withdrew from Pyxis two (2) 5/500mg tablets of Hydrocodone and charted on the MAR that the medication was administered at 1623 hours. The pain assessment was not charted.
- e. Summary: No pain assessments were charted for the 1245 and 1623 administrations.

22 Patient FIN 9024432:

- a. The physician's order(s) for this patient were as follows: Lorazepam 0.5mg every four (4) hours as needed for anxiety, and Vicodin 5/500mg, 1-2 tablets every four (4) hours as needed for moderate/severe pain (4-10 of 10).
- b. On February 12, 2009, at 0757 hours, Respondent withdrew from Pyxis one (1) 0.5mg tablet of Lorazepam and charted on the MAR that the medication was she administered 0.5mg Ativan at 0813 hours. No issue.
- c. On February 12, 2009, at 0830 hours, Respondent charted on the MAR that she administered 0.5mg Ativan to the patient. There is no record of removal of Ativan from Pyxis.

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- d. On February 12, 2009, at 0832 hours, Respondent withdrew from Pyxis one (1) 5/500mg tablet Hydrocodone and charted on the MAR that the medication was administered at 0845 hours. The pain assessment charted at "0" at 0700 hours.
- e. On February 12, 2009, at 0914 hours, Respondent withdrew from Pyxis one (1) 5/500mg tablet of Hydrocodone and charted on the MAR that the medication was administered at 0900 hours. The pain assessment was charted as "0" at 0800 hours and 1200 hours and the medication was charted as given 14 minutes before removal and given more frequently than every four (4) hours.
- f. On February 12, 2009, at 1143 hours, Respondent withdrew from Pyxis two (2) 5/500mg tablet of Hydrocodone and charted on the MAR that the medication was administered at 1230 hours. The pain assessment was charted as "0" at 0800 hours and 1200 hours and the medication was charted as given 47 minutes after removal and given more frequently than every four (4) hours.
- g. On February 12, 2009, at 1531 hours, Respondent withdrew from Pyxis two (2) 5/500mg tablets of Hydrocodone and charted on the MAR as having been given at 1545 hours. The pain level charted as "3" at 1500 hours and given more frequently than every four (4) hours.
- h. On February 12, 2009, at 1801 hours, Respondent withdrew from Pyxis one (1) 0.5mg tablet Lorazepam and charted on the MAR that the medication was administered at 1749 hours. The medication was charted as given 12 minutes before removal.
- i. On February 12, 2009, at 1848 hours, Respondent withdrew from Pyxis one (2) 5/500 tablets of Hydrocodone and charged on the MAR that the medication was administered at 1914 hours.
- j. Respondent removed eight (8) 5/500mg tablets of Hydrocodone, eight (8) tablets are charted as given with pain level less than "4", one (1) tablet charted as given prior to removal, two (2) tablets charted as given 47 minutes after removal. Respondent also removed two (2) tablets of Lorazepam and charted the administering of three (3) 0.5mg tablets and one (1) 0.5mg tablet was charted as given 12 minutes before it was removed.

23. Patient FIN 9024476:

- a. The physician's order(s) for this patient were as follows: Vicodin 5/500mg tablet every four (4) hours as needed for severe pain.
- b. On February 12, 2009, at 1218 hours, Respondent withdrew from Pyxis one (1) 5/500mg tablet of Hydrocodone charted on the MAR that she administered the medication at 1215 hours. Pain assessment was charted as 7, and the administration of the medication as charted as having been given three minutes after removal.
- c. On February 12, 2009, at 1631 hours, Respondent withdrew from Pyxis one (1) 5/500mg tablet of Hydrocodone charted on the MAR that she administered the medication at 1630 hours. Pain assessment was charted as 6 by someone other than Respondent.
- d. On February 12, 2009, at 1857 hours, Respondent withdrew from Pyxis one (1) 5/500mg tablet of Hydrocodone charted on the MAR that she administered the medication at 1915 hours. Pain assessment was charted as 9, and the medication was administered more frequently than every 4 hours.
- e. Summary: The time between the second and third dose is 2 hours and 45 minutes.

24. Patient FIN 9025586:

- a. The physician's order(s) for this patient were as follows: Lorazepam 0.5mg every six (6) hours as needed for anxiety.
- b. On February 16, 2009, at 0820 hours, Respondent withdrew from Pyxis one (1) 0.5mg tablet of Lorazepam and charted on the MAR that she administered the medication at 0845 hours.
- c. On February 16, 2009, at 1406 hours, Respondent withdrew from Pyxis one (1) 0.5mg tablet of Lorazepam and charted on the MAR that she administered the medication at 1420 hours.
- d. On February 16, 2009, at 1637 hours, Respondent withdrew from Pyxis one (1) 0.5mg tablet of Lorazepam and "wasted" it at 1819 hours. The wastage was witnessed by witness ID #22849.

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25. Patient FIN 9024043:

- a. The physician's order(s) for this patient were as follows: Hydrocodone 5/500mg, 2 tablets every six (6) hours as needed.
- b. On February 20, 2009, at 0807 hours, Respondent withdrew from Pyxis two (2) 5/500mg tablet of Hydrocodone and charted on the MAR that she administered the medication at 0800 hours. The pain assessment was charted as "0" and the medication given before removed.
- c. Summary: The medication was charted as given to the patient with "0" level of pain.

26. Patient FIN 9027194:

- a. The physician's order(s) for this patient were as follows: Hydrocodone 5/500mg, 1-2 tablets every four (4) hours as needed for severe to moderate pain.
- b. On February 20, 2009, at 0847 hours, Respondent withdrew from Pyxis two (2) 5/500mg tablet of Hydrocodone and charted on the MAR that she administered the medication at 0900 hours. The pain assessment was charted as "0."
- c. On February 20, 2009, at 1344 hours, Respondent withdrew from Pyxis two (2) 5/500mg tablet of Hydrocodone and charted on the MAR that she administered the medication at 1340 hours. The pain assessment was charted as "0."

27. Patient FIN 9031718:

- a. The physician's order(s) for this patient were as follows: Lorazepam 0.5 mg. every four (4) hours as needed for anxiety; Hydrocodone 5/500mg, 1 (one) tablet every four (4) hours as needed.
- b. On February 26, 2009, at 0831 hours, Respondent withdrew from Pyxis one (1) 0.5mg Lorazepam and charted on the MAR that she administered the medication at 0941 hours, seventy (70) minutes after removal.
- c. On February 26, 2009, at 1114 hours, Respondent withdrew from Pyxis one (1) 5/500mg tablet of Hydrocodone and charted on the MAR that she administered the medication at 1126 hours. The pain assessment was charted as "5" at 1100 hours.

- d. On February 26, 2009, at 1249 hours, Respondent withdrew from Pyxis one (1) 0.5mg tablet of Lorazepam and charted on the MAR that she administered the medication at 1300 hours. This medication was given more frequently than every four (4) hours by 41 minutes.
- e. On February 26, 2009, at 1600 hours, Respondent withdrew from Pyxis one (1) 5/500mg tablet of Hydrocodone and charted on the MAR that she administered the medication at 1606 hours. There was no pain assessment charted.
- f. On February 26, 2009, at 1728 hours, Respondent withdrew from Pyxis one (1) 0.5mg tablet of Lorazepam and charted on the MAR that she administered the medication at 1900 hours. This medication was charted as given 1 hour 32 minutes after removal.
- g. On February 26, 2009, at 1914 hours, Respondent withdrew from Pyxis one (1) 5/500mg tablet of Hydrocodone. This medication was not charted as given to the patient and there was no pain assessment charted. One (1) tablet of Hydrocodone is unaccounted for.
- h. On February 26, 2009, at 1915 hours, Respondent withdrew from Pyxis one (1) 0.5mg tablet of Lorazepam. This medication was not charted as given to the patient. One (1) 0.5mg tablet of Lorazepam is unaccounted for.
- i. Summary: Respondent removed three 5/500mg tablets of Hydrocodone, two (2) were charted as given and one (1) is unaccounted for. Respondent removed four (4) 0.5mg tablets of Lorazepam, three (3) were charted as given and one (1) is unaccounted for. On one occasion, Respondent charted giving one (1) 0.5mg tablet 71 minutes after it was removed, on a second occasion, Respondent charted giving one (1) 0.5mg tablet 1 hour 32 minutes after it was removed.

28. Patient FIN 9032615:

- a. The physician's order(s) for this patient were as follows: Vicodin 5/500mg 1-2 tablets every four (4) hours as needed for moderate to severe pain (4-10).
- b. On February 26, 2009, at 0801 hours, Respondent withdrew from Pyxis two (2) 5/500mg tablets of Hydrocodone and charted on the MAR that she administered one (1) tablet at 0830 hours. The pain assessment is unreadable. One (1) 5/500mg tablet of Hydrocodone is unaccounted for.

- c. On February 26, 2009, at 1130 hours, Respondent withdrew from Pyxis two (2) 5/500mg tablets of Hydrocodone and charted on the MAR that she administered two (2) tablets at 1200 hours. There was no pain assessment charted and the medication was charted as given more frequently than every four hours (3 hrs. 30 minutes).
- d. On February 26, 2009, at 1422 hours, Respondent withdrew from Pyxis two (2) 5/500mg Hydrocodone and charted on the MAR that she administered two (2) tablet at 1500 hours. No pain assessment was charted and the medication was charted as given more frequently than every four hours (3 hours).
- e. Summary: Respondent removed six (6) 5/500mg tablets of Hydrocodone, five (5) tablets were charted as given, and one (1) tablet is unaccounted for. The second and third doses were given less than every four (4) hours as ordered by the physician. The pain assessments were unreadable or not charted.

29. Patient FIN 9033252:

- a. The physician's order(s) for this patient were as follows: Vicodin 1-2 5/500mg tablets every four (4) hours for moderate to severe pain; Lorazepam 0.5 mg. every four (4) hours as needed for anxiety.
- b. On February 26, 2009, at 1041 hours, Respondent withdrew from Pyxis two (2) 5/500 tablets of Hydrocodone and charted on the MAR that she administered the medication at 1106 hours. The medication was charted as given when the pain level was at "0."
- c. On February 26, 2009, at 1041 hours, Respondent withdrew from Pyxis one (1) 0.5mg tablet of Lorazepam and charted on the MAR that she administered the medication at 1107 hours.
- d. On February 26, 2009, at 1558 hours, Respondent withdrew from Pyxis two (2) 5/500 tablets of Hydrocodone and charted on the MAR that she administered the medication at 1607 hours. The medication was charted as given when the pain level was at "0."
- e. On February 26, 2009, at 1558 hours, Respondent withdrew from Pyxis one (1) 0.5mg tablet of Lorazepam and charted on the MAR that she administered the medication at 1607 hours.

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- f. On February 26, 2009, at 1558 hours, Respondent withdrew from Pyxis two (2) 5/500 tablets of Hydrocodone and charted on the MAR that she administered the medication at 1607 hours. The medication was charted as given when the pain level was at "0."
- g. On February 26, 2009, at 1856 hours, Respondent withdrew from Pyxis two (2) 5/500 tablets of Hydrocodone and charted on the MAR that she administered the medication at 1945 hours. The medication was charted as given when the pain level charted was at "0," charted as given more frequently than every four (4) hours (3 hours) as ordered by the physician, and charted as given 49 minutes after removal.
- h. On February 26, 2009, at 1857 hours, Respondent withdrew from Pyxis one (1) 0.5mg tablet of Lorazepam and charted on the MAR that she administered the medication at 1946 hours. The medication was charted as given more frequently than every four (4) hours (3 hours), and charted as given 49 minutes after removal.
- i. Summary: Respondent removed six (6) 5/500mg tablets of Hydrocodone, six (6) tablets were charted as given with "0" pain levels. The time between the second and third doses were given less than every four (4) hours as ordered by the physician. Respondent removed three (3) 0.5mg tablets of Lorazepam, three (3) tablets were charted as given and the time between the second and third does were given less than every four hours as ordered by the physician.

30. Patient FIN 9032866:

- a. The physician's order(s) for this patient were as follows: Lorazepam 0.5 mg. every four (4) hours as needed for anxiety.
- b. On February 27, 2009, at 1059 hours, Respondent withdrew from Pyxis one (1) 0.5mg tablet of Lorazepam and charted on the MAR that she administered the medication at 1107 hours.
- c. On February 26, 2009, at 1302 hours, Respondent "wasted" one (1) 0.5mg Lorazepam tablet, witnessed by CMG8. This was possibly the tablet withdrawn as described in paragraph 28(b).

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- d. On February 27, 2009, at 1318 hours, Respondent withdrew from Pyxis one (1) 0.5mg tablet of Lorazepam and not charted as having been given to the patient. One (1) 0.5mg tablet of Lorazepam is unaccounted for.
- c. On February 27, 2009, at 1636 hours, Respondent withdrew from Pyxis one (1) 0.5mg tablet of Lorazepam and charted on the MAR that she administered the medication at 1643 hours.
- f. On February 27, 2009, at 1856 hours, Respondent withdrew from Pyxis one (1) 0.5mg tablet of Lorazepam and not charted as having been to the patient. One (1) 0.5mg tablet of Lorazepam is unaccounted for.
- g. Summary: Respondent removed four (4) 0.5mg tablets of Lorazepam, two (2) tablets are charted as given, two (2) tablets are unaccounted for, and one (1) tablet wasted.

31. Patient FIN 9032925:

- a. The physician's order(s) for this patient were as follows: Lorazepam 0.5 mg. every four (4) hours as needed for anxiety.
- b. On February 27, 2009, at 1722 hours, Respondent withdrew from Pyxis one (I) 0.5mg tablet of Lorazepam and charted on the MAR that she administered the medication at 1845 hours (1 hour and 23 minutes after removal).
- c. On February 27, 2009, at 1859 hours, Respondent withdrew from Pyxis one (1) 0.5mg tablet of Lorazepam and not charted as having been given to the patient. There is one (1) 0.5mg tablet Lorazepam unaccounted for.
- d. Summary: Respondent removed two (2) 0.5mg tablets of Lorazepam, one (1) is charted as given 1 hour and 23 minutes after being removed, and one (1) is unaccounted for.

32. Patient FIN 9026860:

- a. The physician's order(s) for this patient were as follows: Vicodin 5/500mg 2 tablets every 4-6 hours as needed for pain.
- b. On February 28, 2009, at 0903 hours, Respondent withdrew from Pyxis two 5/500mg tablets of Hydrocodone and charted on the MAR that she administered the medication at 0911 hours. The medication was given with a charted "0" pain level.

c. On February 28, 2009, at 1128 hours, Respondent withdrew from Pyxis two 5/500mg tablets of Hydrocodone and charted on the MAR that she administered the medication at 1200 hours. The medication was given with a charted "0" pain level and removed and given 2 hours 49 minutes after the previous dose, not every 4-6 hours as needed for pain per the physician's orders.

- d. On February 28, 2009, at 1517 hours, Respondent withdrew from Pyxis two 5/500mg tablets of Hydrocodone. The medication was not charged as having been given. Two (2) 5/500 tablets of Hydrocodone are unaccounted for.
- e. Summary: Respondent removed six (6) 5/500 mg. tablets of Hydrocodone, four (4) are charted as given with "0" pain level, and two (2) are unaccounted for. The second dose was given 2 hours 49 minutes after the previous dose, not every 4-6 hours as needed for pain, as ordered by the physician.

33. Patient FIN 9032256:

- a. The physician's order(s) for this patient were as follows: Vicodin 5/500mg 1 tablet every 4 hours as needed for pain.
- b. On February 28, 2009, at 0925 hours, Respondent withdrew from Pyxis one 5/500mg tablets of Hydrocodone and charted on the MAR that she administered the medication at 1014 hours. The medication was charted as given with no pain assessment and removed 49 minutes before given.
- c. On February 28, 2009, at 1141 hours, Respondent withdrew from Pyxis one 5/500mg tablets of Hydrocodone and charted on the MAR that she administered the medication at 1148 hours. The medication was charted as given with "0" pain level, and removed and given 1 hour 34 minutes after the previous dose, not every four hours as needed for pain per the physician's orders.
- d. Summary: On November 28, 2009, Respondent removed two (2) 5/500 tablets of Hydrocodone, two (2) were charted as given with "0" pain level and second tablet was administered 1 hour 34 minutes after the first dose given.

FIRST CAUSE FOR DISCIPLINE

(False Entries in Hospital/Patient Records)

34. Respondent is subject to disciplinary action under section 2761(a), on the grounds of unprofessional conduct, as defined in Code section 2762(e), in that between or about February 3, 2009 and February 28, 2009, while on duty as a registered nurse at Palomar Medical Center, Escondido, California, Respondent falsified, or made grossly incorrect, grossly inconsistent, or unintelligible entries in hospital, patient, or other records pertaining to the controlled substances Hydrocodone and Lorazepam, as is more fully detailed in paragraphs 15 through 33, above, which are incorporated herein by reference.

SECOND CAUSE FOR DISCIPLINE

(Obtain or Possess Controlled Substances)

35. Respondent is subject to disciplinary action pursuant to Code section 2761(a), on the grounds of unprofessional conduct, as defined by Code section 2762(a), in that between or about February 3, 2009 and February 28, 2009, while on duty as a registered nurse at Palomar Medical Center, Escondido, California, Respondent obtained and possessed the controlled substances Hydrocodone and Lorazepam, in violation of Code section 4060 and Health and Safety Code sections 11170 and 11173, as is set forth in paragraphs 15 through 33, above, which are incorporated herein by reference.

PRAYER

WHEREFORE, Complainant requests that a hearing be held on the matters herein alleged, and that following the hearing, the Board of Registered Nursing issue a decision:

- 1. Revoking or suspending Registered Nurse License Number 748793, issued to Christine Marie Nowicki;
- 2. Ordering Christine Marie Nowicki to pay the Board of Registered Nursing the reasonable costs of the investigation and enforcement of this case, pursuant to Business and Professions Code section 125.3;

1	3. Taking such other and further action as deemed necessary and proper.								
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